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(54) Title: POLYMERIZATION OF ETHYLENE		
(57) Abstract		
Ethylene is polymerized by novel nickel or iron complexes of selected novel 2,2-bipyridine-6-imines, optionally with the addition of other cocatalysts. The polymers are useful for films and molding resins.		

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TITLE

## POLYMERIZATION OF ETHYLENE

FIELD OF THE INVENTION

5       Ethylene is polymerized using selected iron or nickel complexes of 2,2'-bipyridine-6-imines, optionally in the presence of certain activators such alkylaluminum compounds. Also disclosed herein are novel 2,2'-bipyridine-6-imines and their iron and nickel complexes.

10

TECHNICAL BACKGROUND

Polyethylene is a very important item of commerce, large quantities of various grades of this polymer being produced annually for a large number of uses, such as packaging films and moldings. There are many different methods for making such polyethylenes, including many used commercially, such as free radical polymerization to make low density polyethylene, and many so-called coordination catalysts such as Ziegler-Natta-type and metallocene-type catalysts. Each of these catalyst systems has its advantages and disadvantages, including cost of the polymerization and the particular structure of the polyethylene produced. Due to the importance of polyethylene, new catalyst systems which are economical and/or produce new types of polyethylenes are constantly being sought.

US5866663 describes the polymerization of ethylene and other olefins using nickel and other transition metal complexes of various imines and similar compounds. This publication is incorporated by reference herein for all purposes.

US5955555, WO98/30612, WO98/38228, WO99/02472 and WO99/12981 (all incorporated by reference herein for all

purposes) describe the use of iron or cobalt complexes of 2,6-diacylpyridinebisimines or 2,6-pyridinedicarboxaldehydebisimines as catalysts for the polymerization of olefins, mostly of ethylene. These publications describe the preparation of polyethylenes ranging in molecular weight from low molecular weight alpha-olefins and other oligomers to high molecular weight polyethylenes.

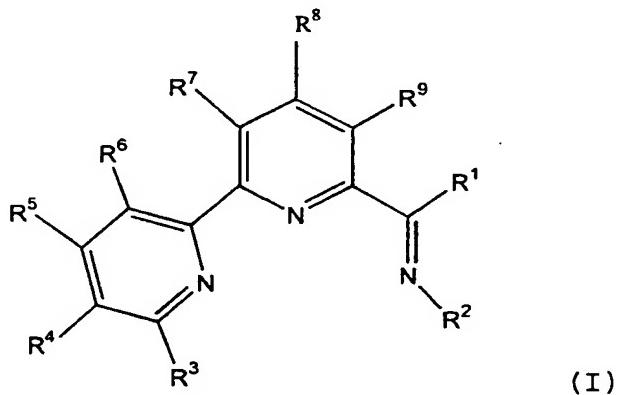
EP-A-0454231 (also incorporated by reference herein for all purposes) describes the use of various complexes of transition metals with selected ligands such as "bipyridine" to polymerize ethylene or other olefins.

In all of the publications cited above, the ligands described herein are not mentioned.

15

#### SUMMARY OF THE INVENTION

This invention concerns, a first process for the production of polyethylene, comprising the step of contacting, at a temperature of about -100°C to about +200°C, ethylene and a Ni or Fe complex of a ligand of the formula



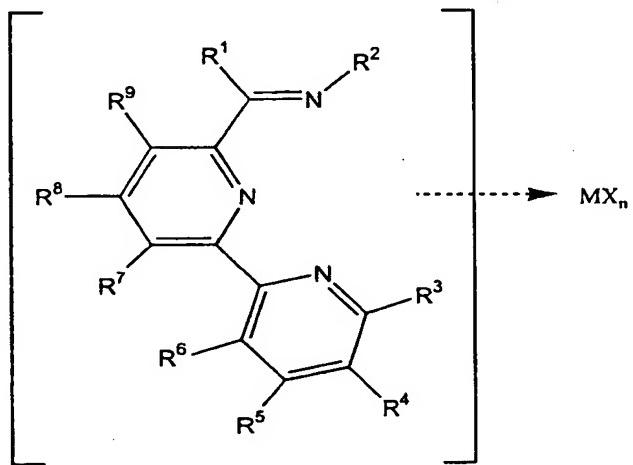
wherein:

R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it; and

each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> vicinal to one another, or R<sup>6</sup> and R<sup>7</sup> taken together, 5 may form a ring.

Also disclosed herein is a second process for the production of polyethylene, comprising the step of contacting, at a temperature of about -100°C to about +200°C, ethylene, a compound of the formula



(II)

10

and:

(a) a first compound W, which is a neutral Lewis acid capable of abstracting X<sup>-</sup> an alkyl group or a hydride group from M to form WX<sup>-</sup>, (WR<sup>20</sup>)<sup>-</sup> or WH<sup>-</sup>, and which is also capable of transferring an alkyl group or a hydride to M, provided that WX<sup>-</sup> is a weakly coordinating anion; or

(b) a combination of second compound which is capable of transferring an alkyl or hydride group to M and a third compound which is a neutral Lewis acid which is capable of abstracting X<sup>-</sup>, a hydride or an alkyl group from M to form a weakly coordinating anion;

wherein:

M is Fe or Ni;

each X is an anion;

n is an integer so that the total number of negative charges on said anion or anions is equal to the oxidation state of M;

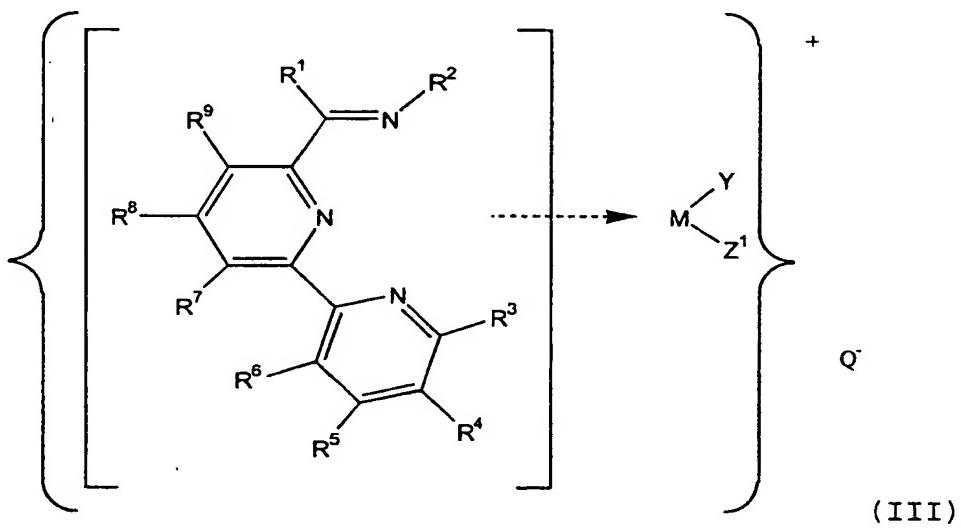
R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

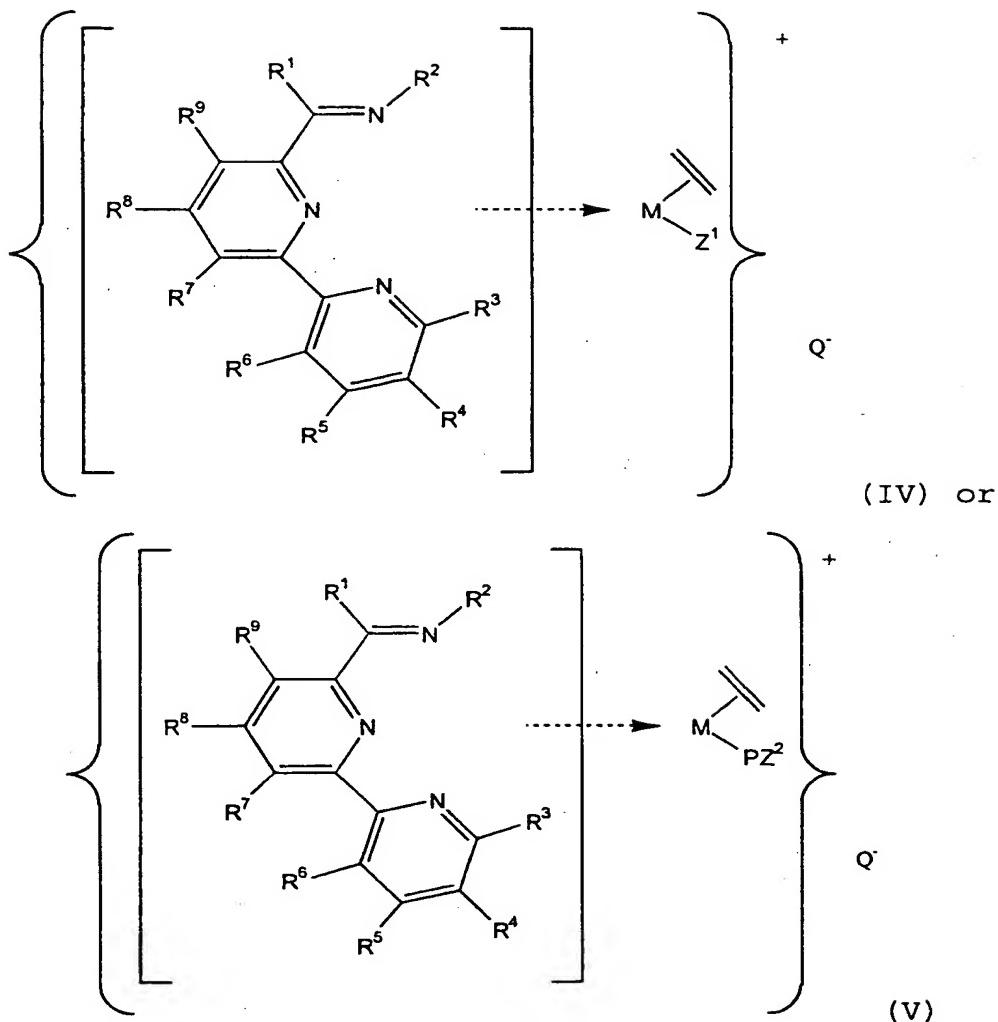
5 R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it; and

each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a

10 functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> vicinal to one another, or R<sup>6</sup> and R<sup>7</sup> taken together, may form a ring.

This invention also concerns a third process for the production of polyethylene, comprising the step of  
15 contacting, at a temperature of about -100°C to about +200°C, ethylene and a compound of the formula





wherein:

M is Ni or Fe;

each X is an anion;

R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it;

each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> vicinal to one another, or R<sup>6</sup> and R<sup>7</sup> taken together, may form a ring;

$Z^1$  is hydride or alkyl or an anionic ligand into which ethylene can insert:

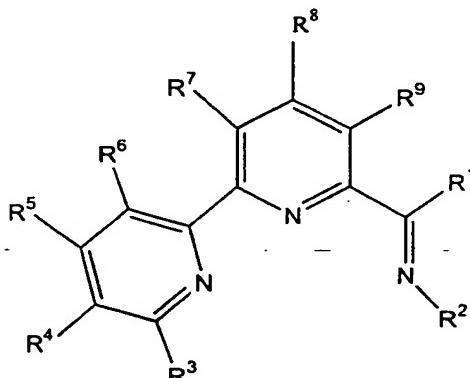
$\text{Y}$  is a neutral ligand capable of being displaced by ethylene, or a vacant coordination site;

5 Q is a relatively non-coordinating anion;

P is a divalent polyethylene group containing one or more ethylene units; and

$Z^2$  is an end group.

Also described herein is a compound of the formula



10

(I)

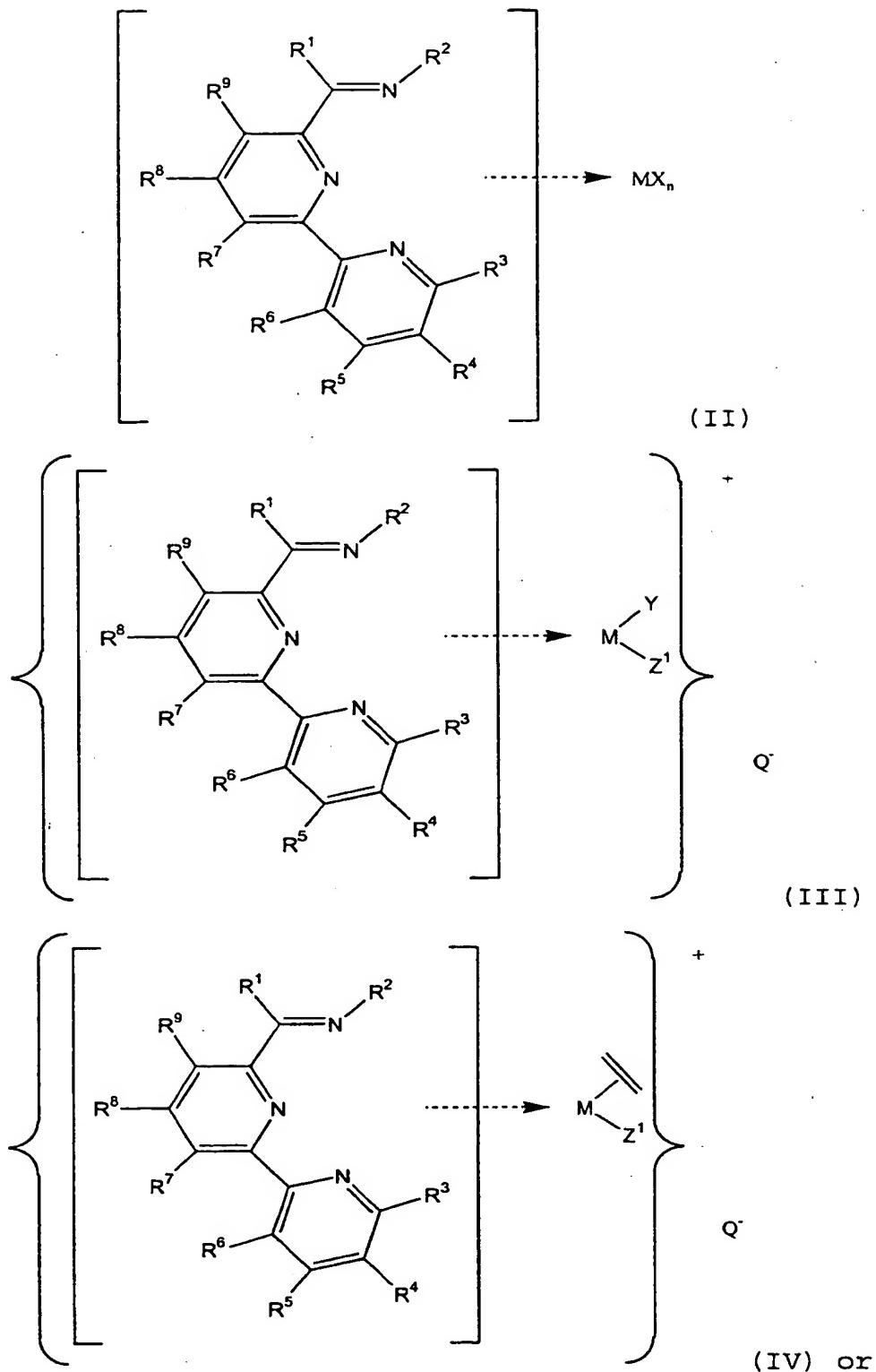
wherein:

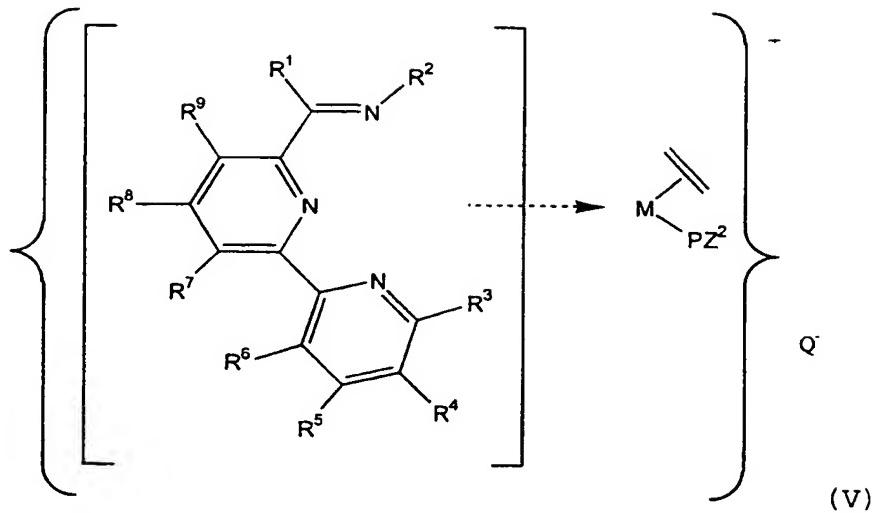
$R^1$  is hydrogen, hydrocarbyl or substituted hydrocarbyl;

$R^2$  is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of  $R^2$  bound to an imino nitrogen atom has at least two carbon atoms bound to it; and

each of  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^8$  and  $R^9$  is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^8$  and  $R^9$  vicinal to one another or,  $R^6$  and  $R^7$  taken together, may form a ring.

Described herein is a compound of the formula





wherein:

M is Fe or Ni;

each X is an anion;

5 n is an integer so that the total number of negative charges on said anion or anions is equal to the oxidation state of M;

R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

10 R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it;

15 each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> vicinal to one another, or R<sup>6</sup> and R<sup>7</sup> taken together, may form a ring;

Z<sup>1</sup> is hydride or alkyl or an anionic ligand into which ethylene can insert;

20 Y is a neutral ligand capable of being displaced by ethylene, or a vacant coordination site;

Q is a relatively non-coordinating anion;

P is a divalent polyethylene group containing one or more ethylene units; and

Z<sup>2</sup> is an end group.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Herein, certain terms are used. Some of them are:

A "hydrocarbyl group" is a univalent group containing only carbon and hydrogen. If not otherwise stated, it is 5 preferred that hydrocarbyl groups herein contain 1 to about 30 carbon atoms.

By "substituted hydrocarbyl" herein is meant a hydrocarbyl group which contains one or more substituent groups which are inert under the process conditions to which 10 the compound containing these groups is subjected. The substituent groups also do not substantially interfere with the process. If not otherwise stated, it is preferred that substituted hydrocarbyl groups herein contain 1 to about 30 carbon atoms. Included in the meaning of "substituted" are 15 heteroaromatic rings.

By "(inert) functional group" herein is meant a group, other than hydrocarbyl or substituted hydrocarbyl, which is inert under the process conditions to which the compound containing the group is subjected. The functional groups 20 also do not substantially interfere with any process described herein that the compound in which they are present may take part in. Examples of functional groups include halo (fluoro, chloro, bromo and iodo), ether such as -OR<sup>18</sup> where R<sup>18</sup> is hydrocarbyl or substituted hydrocarbyl. In 25 cases in which the functional group may be near a nickel or iron atom, the functional group should not coordinate to the metal atom more strongly than groups in (I) which are shown as coordinating to the metal atom in the complexes, that is they should not displace the desired coordinating group.

30 By an "alkyl aluminum compound" is meant a compound in which at least one alkyl group is bound to an aluminum atom. Other groups such as alkoxide, hydride and halogen may also be bound to aluminum atoms in the compound.

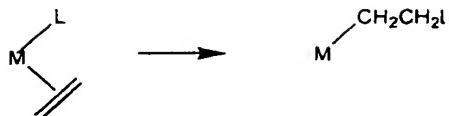
By "neutral Lewis base" is meant a compound, which is not an ion, that can act as a Lewis base. Examples of such compounds include ethers, amines, sulfides, and organic nitriles.

5 By "cationic Lewis acid" is meant a cation that can act as a Lewis acid. Examples of such cations are sodium and silver cations.

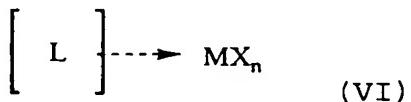
By "relatively noncoordinating anions" (or "weakly coordinating anions" is meant those anions as are generally referred to in the art in this manner, and the coordinating ability of such anions is known and has been discussed in the literature, see for instance W. Beck., et al., Chem. Rev., vol. 88 p. 1405-1421 (1988), and S. H. Stares, Chem. Rev., vol. 93, p. 927-942 (1993), both of which are hereby included by reference. Among such anions are those formed from the aluminum compounds in the immediately preceding paragraph and  $X^-$ , including  $R^9_3AlX^-$ ,  $R^9_2AlClX^-$ ,  $R^9AlCl_2X^-$ , and " $R^9AlOX^-$ ", wherein  $R^9$  is alkyl. Other useful noncoordinating anions include  $BAF^-$  { $BAF$  = tetrakis[3,5-bis(trifluoromethyl)phenyl]borate},  $SbF_6^-$ ,  $PF_6^-$ , and  $BF_4^-$ , trifluoromethanesulfonate, p-toluenesulfonate,  $(R_fSO_2)_2N^-$ , and  $(C_6F_5)_4B^-$ .

By an "empty coordination site" is meant a potential coordination site that is not occupied by a ligand. Thus if an ethylene molecule is in the proximity of the empty coordination site, the ethylene molecule may coordinate to the metal atom.

By a "ligand that may add to ethylene" is meant a ligand coordinated to a metal atom into which an ethylene molecule (or a coordinated ethylene molecule) may insert to start or continue a polymerization. For instance, this may take the form of the reaction (wherein L is a ligand):



Herein a compound of the "formula"



signifies a ligand L [such as (I)] coordinated to the  
5 transition metal group  $\text{MX}_n$ . In this representation whether L  
is acting as a monodentate, bidentate, tridentate, etc.  
ligand is unspecified.

In (I) and its various Ni and Fe complexes as described  
herein it is preferred that:

10  $\text{R}^1$  is hydrogen or hydrocarbyl, more preferably hydrogen  
or alkyl containing 1 to 10 carbon atoms, especially  
preferably hydrogen or methyl; and/or

15  $\text{R}^2$  is aryl or substituted aryl, more preferably phenyl  
or substituted phenyl, especially phenyl substituted in the  
2, or 2 and 6 positions, with alkyl containing 1 to 10  
carbon atoms or halogen, and optionally substituted with  
alkyl containing 1 to 10 carbon atoms in the 4 position;  
and/or

each of  $\text{R}^3$ ,  $\text{R}^4$ ,  $\text{R}^5$ ,  $\text{R}^6$ ,  $\text{R}^7$ ,  $\text{R}^8$  and  $\text{R}^9$  is hydrogen.

20 In specific preferred compounds (I) and complexes  
thereof the following groups are present. In all instances  
each of  $\text{R}^3$ ,  $\text{R}^4$ ,  $\text{R}^5$ ,  $\text{R}^6$ ,  $\text{R}^7$ ,  $\text{R}^8$  and  $\text{R}^9$  is hydrogen:

$\text{R}^1$  is hydrogen and  $\text{R}^2$  is 2,6-diisopropylphenyl; or

$\text{R}^1$  is hydrogen and  $\text{R}^2$  is 2,4,6-tri-t-butylphenyl; or

25  $\text{R}^1$  is hydrogen and  $\text{R}^2$  is pentafluorophenyl; or

$\text{R}^1$  is methyl and  $\text{R}^2$  is 2,6-diisopropylphenyl.

(I) may be prepared by the well-known general reaction  
of a primary amine with a carbonyl compound, in this case a  
2,2'-bipyridine-6-carboxaldehyde or a 6-acyl-2,2'-

bipyridine, to make an imine. Such preparations are illustrated herein in Examples 1-4.

In (II) it is preferred that X is chloride, bromide or acetylacetone (acac).

5        Useful neutral monodentate ligands Y include ethers such as diethyl ether and tetrahydrofuran and nitriles such as acetonitrile.

In the second polymerization process described herein an iron or nickel complex (II) is contacted with ethylene 10 and a neutral Lewis acid W capable of abstracting X<sup>-</sup>, hydride or alkyl from (II) to form a weakly coordinating anion, and must alkylate or be capable of adding a hydride ion to the metal atom, or an additional alkylating agent or an agent capable of adding a hydride anion to the metal atom must be 15 present. The neutral Lewis acid is originally uncharged (i.e., not ionic). Suitable neutral Lewis acids include SbF<sub>5</sub>, Ar<sub>3</sub>B (wherein Ar is aryl) and BF<sub>3</sub>. Suitable cationic Lewis acids or Bronsted acids include NaBAF, silver trifluoromethanesulfonate, HBF<sub>4</sub>, or [C<sub>6</sub>H<sub>5</sub>N(CH<sub>3</sub>)<sub>2</sub>]<sup>+</sup> [B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>]<sup>-</sup>. 20 In those instances in which (II) (and similar catalysts which require the presence of a neutral Lewis acid or a cationic Lewis or Bronsted acid) does not contain an alkyl or hydride group already bonded to the metal atom, the neutral Lewis acid or a cationic Lewis or Bronsted acid also 25 alkylates or adds a hydride to the metal or a separate alkylating or hydriding agent is present, i.e., causes an alkyl group or hydride to become bonded to the metal atom. These alkyl groups, especially methyl, and hydride are groups that may also be Z<sup>1</sup> in (III) and (IV).

30        It is preferred that the alkyl groups in an alkyl aluminum compound contain 1 to 4 carbon atoms each, and more preferred that the alkyl groups are methyl or ethyl.

For instance, alkyl aluminum compounds (see next paragraph) may alkylate (II). However, not all alkyl aluminum compounds may be strong enough Lewis acids to abstract X<sup>-</sup> or an alkyl group from the metal atom. In that case a separate Lewis acid strong enough to do the abstraction must be present.

A preferred neutral Lewis acid, which can alkylate the metal, is a selected alkyl aluminum compound, such as R<sup>19</sup><sub>3</sub>Al, R<sup>19</sup>AlCl<sub>2</sub>, R<sup>19</sup><sub>2</sub>AlCl, and "R<sup>19</sup>AlO" (alkylaluminoxanes), wherein 10 R<sup>19</sup> is alkyl containing 1 to 25 carbon atoms, preferably 1 to 4 carbon atoms. Suitable alkyl aluminum compounds include methylaluminoxane (which is an oligomer with the general formula [MeAlO]<sub>n</sub>), (C<sub>2</sub>H<sub>5</sub>)<sub>2</sub>AlCl, C<sub>2</sub>H<sub>5</sub>AlCl<sub>2</sub>, and [(CH<sub>3</sub>)<sub>2</sub>CHCH<sub>2</sub>]<sub>3</sub>Al.

Metal hydrides such as NaBH<sub>4</sub> may be used to bond hydride groups to the metal M.

In the third polymerization process described herein a nickel or iron complex of (I) is either added to the polymerization process or formed in situ in the process. In fact, more than one such complex may be formed during the 20 course of the process, for instance formation of an initial complex and then reaction of that complex to form a living ended polymer containing such a complex.

Examples of such complexes which may be formed initially in situ include (III) and (IV), as defined above, 25 and wherein the "parallel lines" are an ethylene molecule coordinated to the metal. These complexes may be added directly to the process or formed in situ. For instance, (III) may be formed by the reaction of (II) with a neutral Lewis acid such as an alkyl aluminum compound. Another 30 method of forming such a complex in situ is adding a suitable iron or nickel compound such nickel (II) acetylacetone, (I) and an alkyl aluminum compound. Other metal salts in which anions similar to acetylacetone are

present, and which may be removed by reaction with the Lewis or Bronsted acid. For instance metal halides and carboxylates (such as acetates) may be used, particularly if they are slightly soluble in the process medium. It is  
5 preferred that these precursor metal salts be at least somewhat soluble in the process medium.

After the ethylene polymerization has started, the complex may be in a form such as (V) as defined above wherein P may be a divalent (poly)ethylene group of the  
10 formula - (CH<sub>2</sub>CH<sub>2</sub>)<sub>x</sub>- wherein x is an integer of 1 or more, and Z<sup>2</sup> is an end group (assuming no branching taking place during polymerization), for example the groups listed for Z<sup>1</sup> above. Those skilled in the art will note that (V) is in essence a polymer containing a so-called living end. It is preferred  
15 that M be in +2 oxidation state in (III), (IV) and (V). Compounds such as (III), (IV) and (V) may or may not be stable away from an environment similar to that of the polymerization process, but they may be detected by NMR spectroscopy, particularly one or both of <sup>1</sup>H and <sup>13</sup>C NMR, and  
20 particularly at lower temperatures. Such techniques, especially for polymerization "intermediates" of these types are known, see for instance US5880241, especially Examples 197-203, which is hereby included by reference.

(III), (IV) and (V) may also be used in the absence of  
25 any "co-catalysts" or "activators" to polymerize ethylene in a third polymerization process. Except for the ingredients in the process, the process conditions for the third process, such as temperature pressure, polymerization medium, etc., may be the same as for the first and second  
30 polymerization processes, and preferred conditions for those processes are also preferred for the third polymerization process.

In all the polymerization processes herein, the temperature at which the polymerization is carried out is about -100°C to about +200°C, preferably about 0°C to about 150°C, more preferably about 25°C to about 100°C. A 5 convenient ethylene pressure to use in the polymerization is about 300 kPa to about 275 MPa, more preferably about 2 MPa to about 100 MPa.

The polymerization processes herein may be run in the presence of various liquids, particularly aprotic organic 10 liquids. The catalyst system, ethylene, and polyethylene may be soluble or insoluble in these liquids, but obviously these liquids should not prevent the polymerization from occurring. Suitable liquids include alkanes, cycloalkanes, selected halogenated hydrocarbons, and aromatic- 15 hydrocarbons. Hydrocarbons are the preferred solvent. Specific useful solvents include hexane, toluene, benzene, chloroform, methylene chloride, 1,2,4-trichlorobenzene, p-xylene, and cyclohexane.

The catalysts herein may be "heterogenized" by coating 20 or otherwise attaching them to solid supports, such as silica or alumina. Where an active catalyst species is formed by reaction with a compound such as an alkylaluminum compound, a support on which the alkylaluminum compound is first coated or otherwise attached is contacted with the 25 nickel compound precursor to form a catalyst system in which the active nickel catalyst is "attached" to the solid support. These supported catalysts may be used in polymerizations in organic liquids, as described in the immediately preceding paragraph. They may also be used in 30 so-called gas phase polymerizations in which the olefin(s) being polymerized are added to the polymerization as gases and no liquid supporting phase is present.

It is known that certain transition metal containing polymerization catalysts including those disclosed herein, are especially useful in varying the branching in polyolefins made with them, see for instance previously incorporated US5714556, US5880241, WO98/30610 and WO98/30609. It is also known that blends of distinct polymers, that vary for instance in the properties listed above, may have advantageous properties compared to "single" polymers. For instance it is known that polymers with broad or bimodal molecular weight distributions may be melt processed (be shaped) more easily than narrower molecular weight distribution polymers. Similarly, thermoplastics such as crystalline polymers may often be toughened by blending with elastomeric polymers.

Therefore, methods of producing polymers which inherently produce polymer blends are useful especially if a later separate (and expensive) polymer mixing step can be avoided. However in such polymerizations one should be aware that two different catalysts may interfere with one another, or interact in such a way as to give a single polymer.

In such a process the catalysts disclosed herein can be termed the first active polymerization catalyst. Monomers useful with these catalysts are those described (and also preferred) above.

A second active polymerization catalyst (and optionally one or more others) is used in conjunction with the first active polymerization catalyst. The second active polymerization catalyst may be another late transition metal catalyst, for example as described in previously incorporated US5714556, US5880241, US5955555, WO98/30610 and 98/30609. Other useful types of catalysts may also be used for the second active polymerization catalyst. For instance

so-called Ziegler-Natta and/or metallocene-type catalysts may also be used. These types of catalysts are well known in the polyolefin field, see for instance Angew. Chem., Int. Ed. Engl., vol. 34, p. 1143-1170 (1995), EP-A-0416815 and US5198401 for information about metallocene-type catalysts, and J. Boor Jr., Ziegler-Natta Catalysts and Polymerizations, Academic Press, New York, 1979 for information about Ziegler-Natta-type catalysts, all of which are hereby included by reference. Many of the useful polymerization conditions for all of these types of catalysts and the first active polymerization catalysts coincide, so conditions for the polymerizations with first and second active polymerization catalysts are easily accessible. Oftentimes the "co-catalyst" or "activator" is needed for metallocene or Ziegler-Natta-type polymerizations. In many instances the same compound, such as an alkylaluminum compound, may be used as an "activator" for some or all of these various polymerization catalysts.

Suitable catalysts for the second polymerization catalyst also include metallocene-type catalysts, as described in US5324800 and EP-A-0129368; particularly advantageous are bridged bis-indenyl metallocenes, for instance as described in US5145819 and EP-A-0485823. Another class of suitable catalysts comprises the well-known constrained geometry catalysts, as described in EP-A-0416815, EP-A-0420436, WP-A-0671404, EP-A-0643066 and WO91/04257. Also the class of transition metal complexes described in WO96/13529 can be used.

In one preferred process described herein the first olefin(s) [ethylene polymerized by the first active polymerization catalyst] and second olefin(s) [the monomer(s) polymerized by the second active polymerization catalyst] are identical, and preferred olefins in such a

process are the same as described immediately above. The second olefins may also be a single olefin or a mixture of olefins to make a copolymer. Again it is preferred that they be identical particularly in a process in which 5 polymerization by the first and second active polymerization catalysts make polymer simultaneously.

In some processes herein the first active polymerization catalyst may polymerize a monomer that may not be polymerized by said second active polymerization 10 catalyst, and/or vice versa. In that instance two chemically distinct polymers may be produced. In another scenario two monomers would be present, with one polymerization catalyst producing a copolymer, and the other polymerization catalyst producing a homopolymer, or two 15 copolymers may be produced which vary in the molar proportion or repeat units from the various monomers. Other analogous combinations will be evident to the artisan.

In another variation of this process one of the polymerization catalysts makes an oligomer of an olefin, 20 preferably ethylene, which oligomer has the formula R<sup>70</sup>CH=CH<sub>2</sub>, wherein R<sup>70</sup> is n-alkyl, preferably with an even number of carbon atoms. The other polymerization catalyst in the process then (co)polymerizes this olefin, either by itself or preferably with at least one other olefin, 25 preferably ethylene, to form a branched polyethylene. Preparation of the oligomer (which is sometimes called an α-olefin) by a second active polymerization-type of catalyst can be found in previously incorporated US5880241, and 30 WO99/02472 (also incorporated by reference herein for all purposes).

Likewise, conditions for such polymerizations, using catalysts of the second active polymerization type, will also be found in the appropriate above mentioned references.

Two chemically different active polymerization catalysts are used in this polymerization process. The first active polymerization catalyst is described in detail above. The second active polymerization catalyst may also 5 meet the limitations of the first active polymerization catalyst, but must be chemically distinct. For instance, it may have a different transition metal present, and/or utilize a different type of ligand and/or the same type of ligand which differs in structure between the first and 10 second active polymerization catalysts. In one preferred process, the ligand type and the metal are the same, but the ligands differ in their substituents.

Included within the definition of two active polymerization catalysts are systems in which a single 15 polymerization catalyst is added together with another ligand, preferably the same type of ligand, which can displace the original ligand coordinated to the metal of the original active polymerization catalyst, to produce in situ two different polymerization catalysts.

The molar ratio of the first active polymerization catalyst to the second active polymerization catalyst used will depend on the ratio of polymer from each catalyst desired, and the relative rate of polymerization of each catalyst under the process conditions. For instance, if one 20 wanted to prepare a "toughened" thermoplastic polyethylene that contained 80% crystalline polyethylene and 20% rubbery polyethylene, and the rates of polymerization of the two catalysts were equal, then one would use a 4:1 molar ratio of the catalyst that gave crystalline polyethylene to the catalyst that gave rubbery polyethylene. More than two 25 active polymerization catalysts may also be used if the desired product is to contain more than two different types of polymer.

The polymers made by the first active polymerization catalyst and the second active polymerization catalyst may be made in sequence, i.e., a polymerization with one (either first or second) of the catalysts followed by a 5 polymerization with the other catalyst, as by using two polymerization vessels in series. However it is preferred to carry out the polymerization using the first and second active polymerization catalysts in the same vessel(s), i.e., simultaneously. This is possible because in most instances 10 the first and second active polymerization catalysts are compatible with each other, and they produce their distinctive polymers in the other catalyst's presence. Any of the processes applicable to the individual catalysts may be used in this polymerization process with 2 or more 15 catalysts, i.e., gas phase, liquid phase, continuous, etc.

The polymers produced by this "mixed catalyst" process may vary in molecular weight and/or molecular weight distribution and/or melting point and/or level of crystallinity, and/or glass transition temperature and/or 20 other factors. For copolymers the polymers may differ in ratios of comonomers if the different polymerization catalysts polymerize the monomers present at different relative rates. The polymers produced are useful as molding and extrusion resins and in films as for packaging. They 25 may have advantages such as improved melt processing, toughness and improved low temperature properties.

Hydrogen may be used to lower the molecular weight of polyethylene produced in the first or second processes. It is preferred that the amount of hydrogen present be about 30 0.01 to about 50 mole percent of the ethylene present, preferably about 1 to about 20 mole percent. The relative concentrations of hydrogen and ethylene may be regulated by their partial pressures.

Included herein within the definitions of all the polymerization processes are mixtures of starting materials that lead to the formation in situ of the transition metal compounds specified in all of the polymerization processes.

5 In the Examples, various abbreviations are used:

acac - acetylacetone

DME - 1,2-dimethoxyethane

DSC - Differential Scanning calorimetry

GPC - Gel Permeation Chromatography

10 Mn - number average molecular weight

Mw - weight average molecular weight

PE - polyethylene

PMAO - methylaluminoxane

Melting point was determined by DSC at a heating rate of 10°C/min. The melting point was taken as the peak of the melting endotherm on the second heat. Usually the DSCs of these polymers exhibited broad melting endotherms.

#### Examples 1-5

##### Preparation of Ligands (I)

20 This procedure illustrates the synthesis of the 2,2'-bipyridinephenylimine ligands. A 100 mL round-bottomed flask was charged with 1.0 g (5.43 mmol) of 2,2'-bipyridine-6-carboxaldehyde. Anhydrous methanol (50 mL) was added forming an orange solution. 2,6-Diisopropylaniline (0.96 g, 5.43 mmol) and 3 drops of formic acid were added and the reaction was stirred overnight. A yellow precipitate began forming after approximately 15 min. The reaction mixture was filtered and the product was washed with methanol and dried in vacuo. A yellow solid (1.22 g, 65%), (I-1), was recovered. <sup>1</sup>H NMR (300 MHz, 20°C, CDCl<sub>3</sub>) δ 8.69-7.12 (m, 10H, bipyridine and aromatic protons); 8.39 (s, 1H, backbone H); 2.99 (septet, 2H, CH(CH<sub>3</sub>)<sub>2</sub>); 1.18 (d, 12H, CH(CH<sub>3</sub>)<sub>2</sub>).

Structures of the various ligands (I) are given in Table 1. In all instances R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> were hydrogen.

Table 1

Example	Ligand	R <sup>1</sup>	R <sup>2</sup>
1	(I-1)	H	2,6-diisopropylphenyl
2	(I-2)	H	2,4,6-tri-t-butylphenyl
3	(I-3)	H	pentafluorophenyl
4	(I-4)	CH <sub>3</sub>	2,6-diisopropylphenyl
5	(I-5)	CH <sub>3</sub>	pentafluorophenyl

5

Examples 6-15Preparation of Fe and Ni Complexes

This is a general procedure for the synthesis of NiBr<sub>2</sub> complexes of 2,2'-bipyridinephenylimines. In a drybox, a round bottom flask was charged with (DME)NiBr<sub>2</sub> (0.225 g, 0.728 mmol). Dichloromethane (10 mL) was added forming a peach suspension. 2,2'-Bipyridinephenylimine (0.728 mmol) was added and the resulting solution was stirred 12 h. The reaction mixture was filtered (sometimes through Celite®) and the solvent was removed *in vacuo*. The product was washed with pentane and dried *in vacuo*.

This is a general procedure for the synthesis of Ni(acac)<sub>2</sub> complexes of 2,2'-bipyridinephenylimines. In a drybox, a round bottom flask was charged with Ni(acac)<sub>2</sub> (0.100 g, 0.28 mmol). Tetrahydrofuran (30 mL) was added forming a green suspension. 2,2'-Bipyridinephenylimine (0.728 mmol) was added and the resulting solution was stirred 12 h. The reaction mixture was filtered through Celite® and the solvent was removed *in vacuo*. The product was washed with pentane and dried *in vacuo*.

This is a general procedure for the synthesis of FeCl<sub>2</sub> complexes of 2,2'-bipyridinephenylimines. In a drybox, a

round bottom flask was charged with 2,2'-bipyridinephenylimine (0.58 mmol). Tetrahydrofuran (30 mL) was added forming a yellow solution. FeCl<sub>2</sub> (0.074 g, 0.58 mmol) was added and the resulting suspension was stirred 12 h. The reaction mixture was filtered and the product was washed with tetrahydrofuran and pentane and dried in vacuo.

For each of the complexes made, yields and other data are given in Table 2.

Table 2

Example	Ligand	Metal Compound	Yield g (%)	Color
6	(I-1)	NiBr <sub>2</sub>	0.25 (61)	brown
7	(I-2)	NiBr <sub>2</sub>	0.23 (75)	orange
8	(I-3)	NiBr <sub>2</sub>	- (75)	royal blue
9	(I-4)	NiBr <sub>2</sub>	0.20 (50)	dark green
10	(I-3)	Ni(acac) <sub>2</sub>	0.083 (50)	forest green
11	(I-5)	Ni(acac) <sub>2</sub>	0.089 (52)	pale green
12	(I-1)	FeCl <sub>2</sub>	0.14 (50)	royal blue
13	(I-3)	FeCl <sub>2</sub>	0.20 (70)	royal blue
14	(I-4)	FeCl <sub>2</sub>	0.17 (64)	royal blue
15	(I-2)	FeCl <sub>2</sub>	- (75)	orange

10

Examples 16-23

Polymerizations with Ethylene

This procedure is general for ethylene polymerization. A reactor was charged with Ni or Fe complex (0.03 mmol). 15 1,2,4-Trichlorobenzene (3 mL) was added and the catalyst solution was placed in a -20°C freezer for 15 min. Another layer of 1,2,4-trichlorobenzene (3 mL) was added and the reactor was again placed in a -20°C freezer for 15 min. This procedure separated the catalyst solution from the 20 co-catalyst through a layer of frozen solvent. PMAO co-catalyst (1 mL) was added and the frozen reactor was brought out of the drybox. The reactor was evacuated, charged with ethylene to 6.9 MPa (1000 psi) and the

polymerization run for 16 h. Polyethylene was recovered by precipitation in methanol, washed with 6 M HCl and methanol and dried under high vacuum overnight. Details of the various polymerizations are given in Table 3. By DSC, in  
5 Example 16 the Mn was 540 and the Mw was 2569, and in Example 21 the Mn was 1676 and the Mw was 80,000.

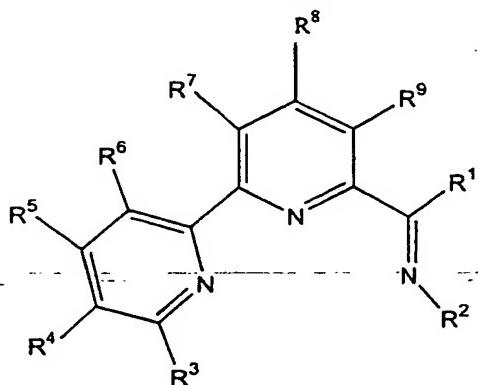
Table 3

Example	Complex of Example	Yield PE mg	Tm, °C
16	11	1200	121
17	14	380	121
18	12	30	124
19	13	50	121
20	5	70	121
21	6	140	128
22	8	20	129
23	10	100	130

CLAIMS

What is claimed is:

1. A process for the production of polyethylene,  
comprising the step of contacting, at a temperature of about  
5 -100°C to about +200°C, ethylene and an active catalyst  
based on a transition metal complex of an imine-functional  
ligand, characterized in that said catalyst is a Ni or Fe  
complex of a ligand of the formula



(I)

10 wherein:

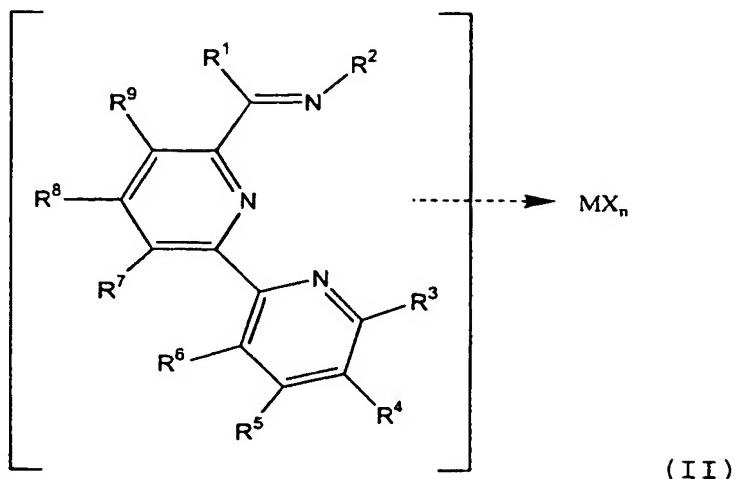
R¹ is hydrogen, hydrocarbyl or substituted hydrocarbyl;

R² is hydrocarbyl or substituted hydrocarbyl, provided  
that a carbon atom of R² bound to an imino nitrogen atom has  
at least two carbon atoms bound to it; and

15 each of R³, R⁴, R⁵, R⁶, R⁷, R⁸ and R⁹ is independently  
hydrogen, hydrocarbyl, substituted hydrocarbyl or a  
functional group, provided that any two of R³, R⁴, R⁵, R⁶, R⁷,  
R⁸ and R⁹ vicinal to one another, or R⁶ and R⁷ taken together,  
may form a ring.

20

2. The process as recited in claim 1, characterized in that said complex is



and also present is:

5           (a) a first compound W, which is a neutral Lewis acid capable of abstracting  $X^-$ , an alkyl group or a hydride group from M to form  $WX^-$ ,  $(WR^{20})^-$  or  $WH^-$ , and which is also capable of transferring an alkyl group or a hydride to M, provided that  $WX^-$  is a weakly coordinating anion; or

10          (b) a combination of second compound which is capable of transferring an alkyl or hydride group to M and a third compound which is a neutral Lewis acid which is capable of abstracting  $X^-$ , a hydride or an alkyl group from M to form a weakly coordinating anion;

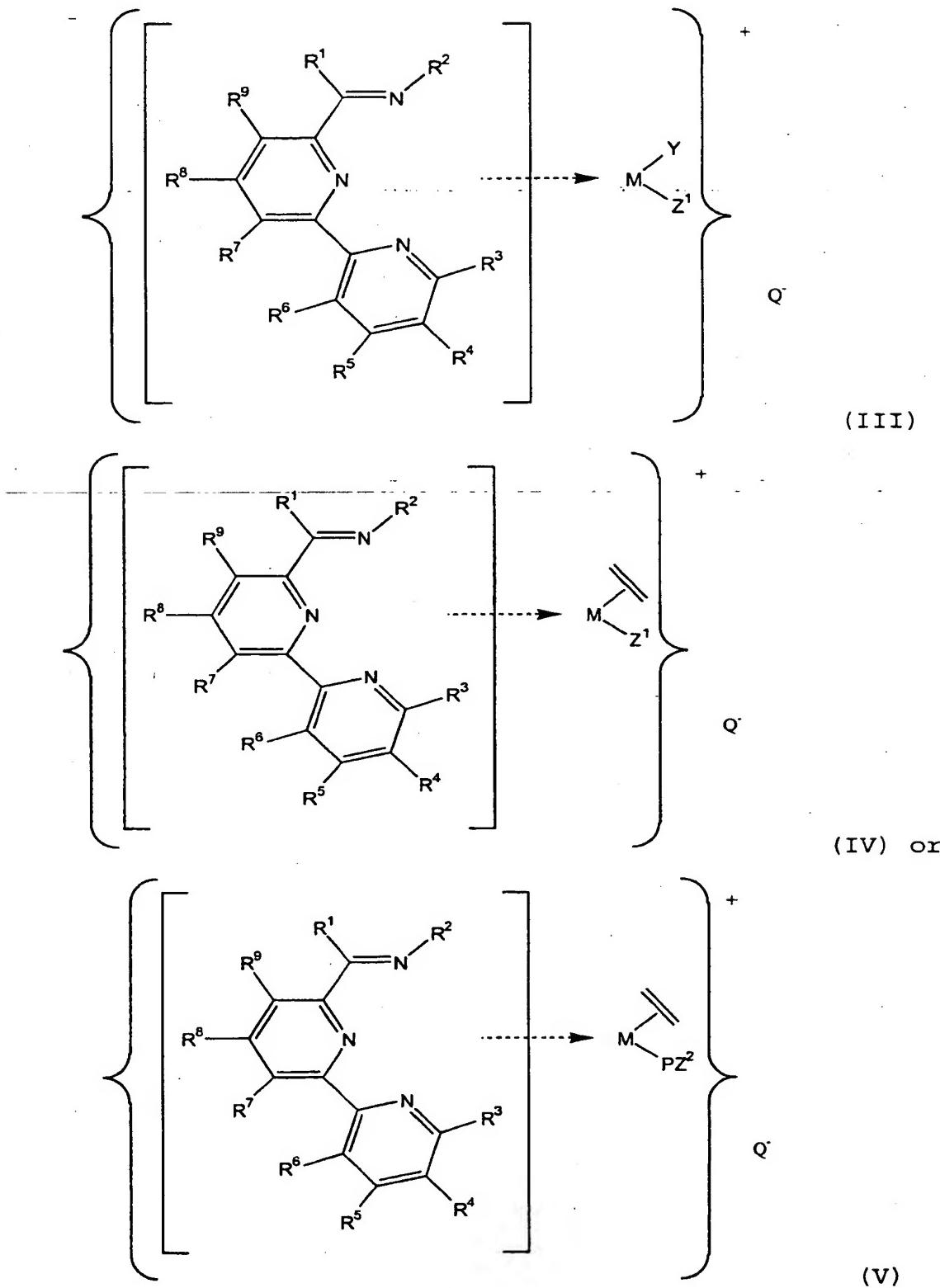
15          wherein:

M is Fe or Ni;

each X is an anion; and

n is an integer so that the total number of negative charges on said anion or anions is equal to the oxidation state of M.

3. The process as recited in claim 1 characterized in that said complex is



wherein:

M is Ni or Fe;

each X is an anion;

5 Z<sup>1</sup> is hydride or alkyl or an anionic ligand into which ethylene can insert;

Y is a neutral ligand capable of being displaced by ethylene or a vacant coordination site;

Q is a relatively non-coordinating anion;

10 P is a divalent polyethylene group containing one or more ethylene molecules; and

Z<sup>2</sup> is an end group.

4. The process as recited in any one of claims 1-3, characterized in that R<sup>1</sup> is hydrogen or alkyl containing 1 to 15 10 carbon atoms, and each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is hydrogen.

5. The process as recited in any one of claims 1-3, characterized in that R<sup>2</sup> is aryl or substituted aryl.

20 6. The process as recited in claim 4, characterized in that R<sup>2</sup> is phenyl or substituted phenyl.

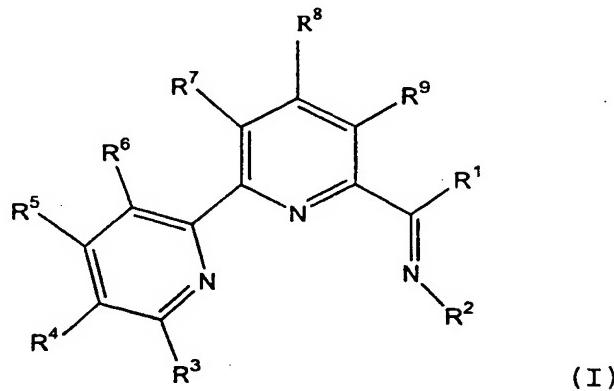
7. The process as recited in any one of claims 1-3, characterize din that said temperature about 25°C to about 25 100°C.

30 8. The process as recited in any one of claims 1-3, characterized in that at least one other polymerization catalyst is present.

9. The process as recited in any one of claims 1-3, characterized in that said process is carried out in the gas phase.

5 10. The process as recited in any one of claims 1-3, characterized in that said process is carried out in the liquid phase.

11. A compound of the formula



10

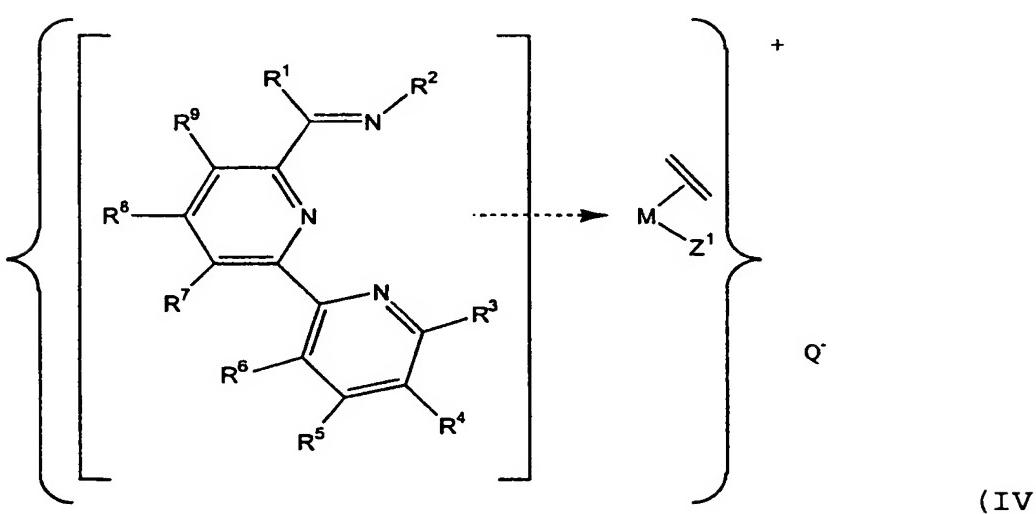
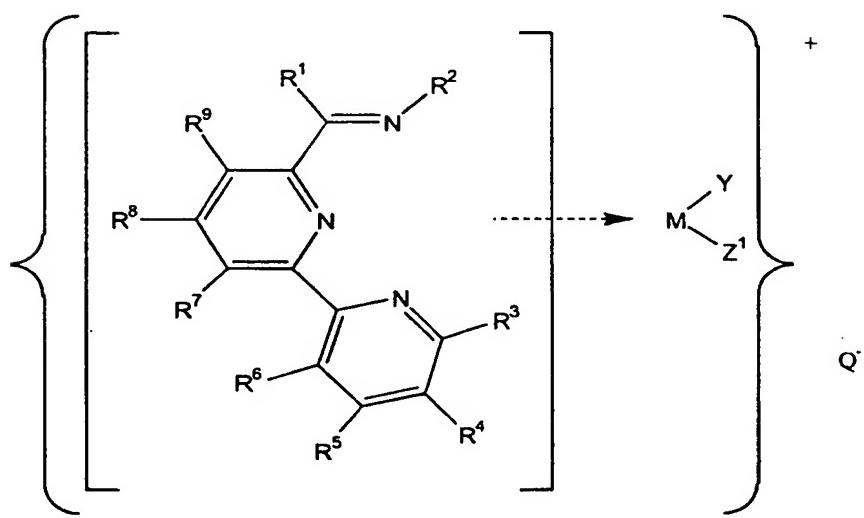
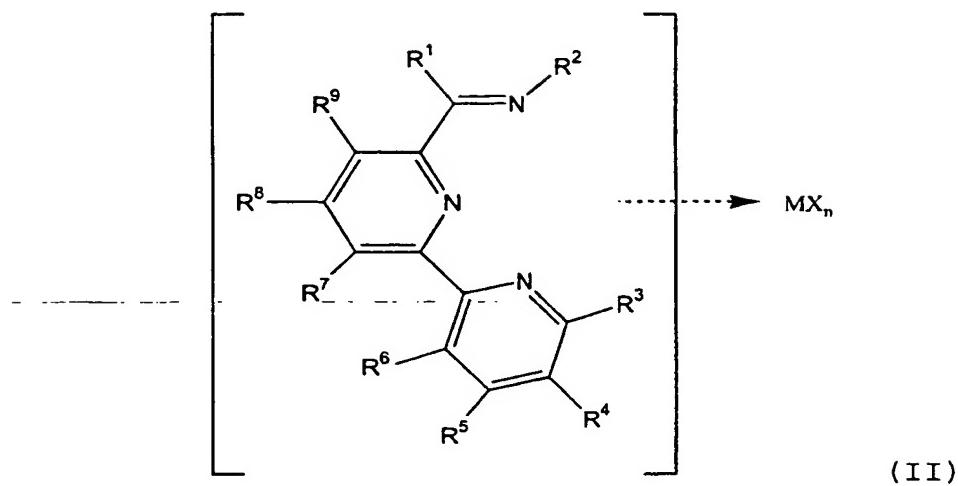
wherein:

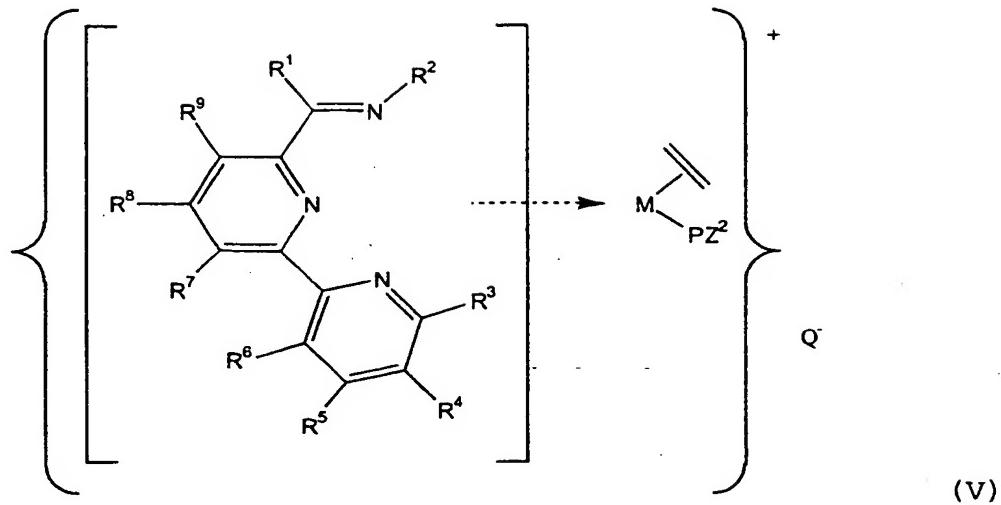
R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it; and

each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> vicinal to one another, or R<sup>6</sup> and R<sup>7</sup> taken together, may form a ring.

## 12. A compound of the formula





wherein:

M is Fe or Ni;

each X is an anion;

5 n is an integer so that the total number of negative charges on said anion or anions is equal to the oxidation state of M;

R<sup>1</sup> is hydrogen, hydrocarbyl or substituted hydrocarbyl;

R<sup>2</sup> is hydrocarbyl or substituted hydrocarbyl, provided

10 that a carbon atom of R<sup>2</sup> bound to an imino nitrogen atom has at least two carbon atoms bound to it;

each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is independently hydrogen, hydrocarbyl, substituted hydrocarbyl or a functional group, provided that any two of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, 15 R<sup>8</sup> and R<sup>9</sup> vicinal to one another or R<sup>6</sup> and R<sup>7</sup> taken together may form a ring;

Z<sup>1</sup> is hydride or alkyl or any other anionic ligand into which ethylene can insert;

Y is a neutral ligand capable of being displaced by ethylene or a vacant coordination site;

Q is a relatively non-coordinating anion;

P is a divalent polyethylene group containing one or more ethylene molecules; and

Z<sup>2</sup> is an end group.

13. The compound as recited in claim 11 or 12,  
characterized in that R<sup>1</sup> is hydrogen or alkyl containing 1 to  
10 carbon atoms, and each of R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is  
5 hydrogen.

14. The compound as recited in claim 11 or 12,  
characterized in that R<sup>2</sup> is aryl or substituted aryl.

10 15. The compound as recited in claim 13, characterized  
in that R<sup>2</sup> is phenyl or substituted phenyl.

16. An olefin polymerization catalyst comprising the  
compound of claim 12.

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 00/11916

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C08F10/00 C08F4/70 C08F2/34 C07D213/53 C07F15/02

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08F C07D C07F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EP0-Internal, WPI Data, CHEM ABS Data

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GOODWIN H A ET AL: "METAL COMPLEXES OF 1,10-PHENANTHROLINE DERIVATIVES. VIII* COMPLEXES OF 1,10-PHENANTHROLINE-2-CARBALDEHYDE IMINES" AUSTRALIAN JOURNAL OF CHEMISTRY, XX, XX, vol. 27, no. 10, 1974, pages 2121-2129, XP000911008 ISSN: 0004-9425 figure 1; table 1	11
A	----- -/-	1-10, 12-16

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

## \* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
6 September 2000	18/09/2000
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3018	Authorized officer  Parry, J

## INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/11916

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; ZIESSEL, RAYMOND ET AL: "Copper(I) helicates containing bridging but non-chelating polypyridine fragments" retrieved from STN Database accession no. 128:56664 XP002146680 abstract &amp; ANGEW. CHEM., INT. ED. ENGL. (1997), 36(22), 2509-2511 ,</p> <p>-----</p>	11,13-15
X	<p>DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; EL-GHAYOURY, ABDELKRIM ET AL: "Cation-induced macroscopic ordering of non-mesomorphic modules - a new application for metallohelicates" retrieved from STN Database accession no. 129:325292 XP002146681 abstract &amp; ANGEW. CHEM., INT. ED. (1998), 37(16), 2205-2208 ,</p> <p>-----</p>	11,13-15
A	<p>DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; KUO, YANG MING: "Electrochemical studies of nickel bis(2,2':6',2''-terpyridine) with alkyl/aryl/allyl bromides and activated olefins in nonaqueous solvents" retrieved from STN Database accession no. 118:134609 XP002146682 abstract &amp; JIEMIAN KEXUE HUIZHI (1992), 15(1), 23-43 ,</p> <p>-----</p>	1-16